



THE UNIVERSITY *of* EDINBURGH

Edinburgh Research Explorer

Deep resequencing unveils genetic architecture of ADIPOQ and identifies a novel low-frequency variant strongly associated with adiponectin variation

Citation for published version:

Warren, LL, Li, L, Nelson, MR, Ehm, MG, Shen, J, Fraser, DJ, Aponte, JL, Nangle, KL, Slater, AJ, Woollard, PM, Hall, MD, Topp, SD, Yuan, X, Cardon, LR, Chissoe, SL, Mooser, V, Morris, AD, Palmer, CNA, Perry, JR, Frayling, TM, Whittaker, JC & Waterworth, DM 2012, 'Deep resequencing unveils genetic architecture of ADIPOQ and identifies a novel low-frequency variant strongly associated with adiponectin variation', *Diabetes*, vol. 61, no. 5, pp. 1297-1301. <https://doi.org/10.2337/db11-0985>

Digital Object Identifier (DOI):

[10.2337/db11-0985](https://doi.org/10.2337/db11-0985)

Link:

[Link to publication record in Edinburgh Research Explorer](#)

Document Version:

Publisher's PDF, also known as Version of record

Published In:

Diabetes

Publisher Rights Statement:

© 2012 by the American Diabetes Association.

Readers may use this article as long as the work is properly cited, the use is educational and not for profit, and the work is not altered. See <http://creativecommons.org/licenses/by-nc-nd/3.0/> for details.

General rights

Copyright for the publications made accessible via the Edinburgh Research Explorer is retained by the author(s) and / or other copyright owners and it is a condition of accessing these publications that users recognise and abide by the legal requirements associated with these rights.

Take down policy

The University of Edinburgh has made every reasonable effort to ensure that Edinburgh Research Explorer content complies with UK legislation. If you believe that the public display of this file breaches copyright please contact openaccess@ed.ac.uk providing details, and we will remove access to the work immediately and investigate your claim.



Deep Resequencing Unveils Genetic Architecture of *ADIPOQ* and Identifies a Novel Low-Frequency Variant Strongly Associated With Adiponectin Variation

Liling L. Warren,¹ Li Li,¹ Matthew R. Nelson,¹ Margaret G. Ehm,¹ Judong Shen,¹ Dana J. Fraser,¹ Jennifer L. Aponte,¹ Keith L. Nangle,¹ Andrew J. Slater,¹ Peter M. Woollard,² Matt D. Hall,² Simon D. Topp,² Xin Yuan,³ Lon R. Cardon,³ Stephanie L. Chisoe,¹ Vincent Mooser,³ Andrew D. Morris,⁴ Colin N.A. Palmer,⁴ John R. Perry,^{5,6} Timothy M. Frayling,⁵ John C. Whittaker,² and Dawn M. Waterworth³

Increased adiponectin levels have been shown to be associated with a lower risk of type 2 diabetes. To understand the relations between genetic variation at the adiponectin-encoding gene, *ADIPOQ*, and adiponectin levels, and subsequently its role in disease, we conducted a deep resequencing experiment of *ADIPOQ* in 14,002 subjects, including 12,514 Europeans, 594 African Americans, and 567 Indian Asians. We identified 296 single nucleotide polymorphisms (SNPs), including 30 amino acid changes, and carried out association analyses in a subset of 3,665 subjects from two independent studies. We confirmed multiple genome-wide association study findings and identified a novel association between a low-frequency SNP (rs17366653) and adiponectin levels ($P = 2.2 \times 10^{-17}$). We show that seven SNPs exert independent effects on adiponectin levels. Together, they explained 6% of adiponectin variation in our samples. We subsequently assessed association between these SNPs and type 2 diabetes in the Genetics of Diabetes Audit and Research in Tayside Scotland (GO-DARTS) study, comprised of 5,145 case and 6,374 control subjects. No evidence of association with type 2 diabetes was found, but we were also unable to exclude the possibility of substantial effects (e.g., odds ratio 95% CI for rs17366653 [0.91–1.58]). Further investigation by large-scale and well-powered Mendelian randomization studies is warranted. *Diabetes* 61:1297–1301, 2012

Adiponectin is an anti-inflammatory adipokine secreted by adipocytes and is inversely associated with the risk of type 2 diabetes (1); however, whether adiponectin is causal or merely a marker of prediabetes is not yet known. Use of genetics through Mendelian randomization (2,3) is one approach to investigate causality; thus the identification of genetic variation affecting adiponectin levels has drawn much attention. Through linkage and association studies, adiponectin levels have been linked to the *ADIPOQ* locus on

chromosome 3q27 (4–8). The majority of adiponectin genetic investigations to date have been limited to common variants, but with the advent of massively parallel sequencing, we can now explore low-frequency variation within this gene as well. Here, we describe results from a deep resequencing experiment of the exons and flanking regions of *ADIPOQ* in 14,002 individuals. We describe the genetic variations observed and report genetic associations with adiponectin levels in a subset of 3,665 individuals with adiponectin measurements. For variants independently associated with adiponectin levels, we further evaluated their impact on type 2 diabetes susceptibility in a cohort of 5,145 type 2 diabetic and 6,374 control subjects.

RESEARCH DESIGN AND METHODS

We sequenced *ADIPOQ* in 14,002 individuals, including 12,514 Europeans, 594 African Americans, and 567 Indian Asians. Adiponectin levels were measured in a subset of 3,665 subjects of European origin from two studies: 1,579 from the Genetic Epidemiology of the Metabolic Syndrome (GEMS) study (9) and 2,086 from the Cohorte Lausannoise (CoLaus) study (10). The GEMS study is a large multinational study designed to explore the genetic basis of the metabolic syndrome. Subjects in our resequencing study were selected based on DNA availability and consisted of 787 dyslipidemic subjects with an elevated plasma triglyceride and a low serum HDL cholesterol and 792 normolipidemic control subjects having the combination of an elevated plasma triglyceride, a low serum HDL cholesterol, and a BMI $>25 \text{ kg/m}^2$. The CoLaus study is a single-center, population-based study to assess the prevalence of cardiovascular risk factors in the population of Lausanne, Switzerland. We included 2,086 subjects in this experiment based on availability of DNA and phenotype assessment. Genotyping was conducted in the Genetics of Diabetes Audit and Research in Tayside Scotland (GO-DARTS) study (11), which includes a total of 12,348 individuals, 5,145 type 2 diabetic subjects and 6,374 normoglycemic, population-based control subjects, all of European U.K. origin.

DNA sequencing and genotyping. All three exons of *ADIPOQ* (NM_004797) plus 50 bases of flanking sequence (NCBI build 36.3) were selected for capture using a custom Roche NimbleGen (Madison, WI) HD2.1M sequence capture array. Paired-end sequencing was conducted for each 48-sample indexed pool. Variants were called using SOAPsnp (12) at a minimum depth of 7 and a minimum consensus quality of 20. Genotyping in the GO-DARTS study was performed using a Kaspar assay (<http://www.kbioscience.co.uk/>).

Adiponectin measurement. Plasma adiponectin levels were measured using the ELISA assay (R&D Systems, Minneapolis, MN).

Statistical methods. Linear regression analyses were carried out in the GEMS and CoLaus studies separately under an additive genetic model adjusted for significant covariates ($P < 0.05$) in each study, including dyslipidemia status, age, sex, collection site, waist and hip circumference in GEMS, and age, sex, waist and hip circumference, BMI, smoking, and alcohol usage in CoLaus. Adiponectin levels were log transformed and the extreme outliers were set to the 99.9 percentile of the distribution. Single nucleotide polymorphisms (SNPs) with at least 10 copies of the minor allele were analyzed individually, whereas nonsynonymous SNPs with <10 copies were aggregated (13). SNPs and subjects with $>20\%$ missing data were excluded from analysis. Multiple

From ¹Quantitative Sciences, GlaxoSmithKline, Research Triangle Park, North Carolina; ²Quantitative Sciences, GlaxoSmithKline, Stevenage, U.K.; ³Quantitative Sciences, GlaxoSmithKline, Upper Merion, Pennsylvania; the ⁴Biomedical Research Institute, Ninewells Hospital and Medical School, University of Dundee, Dundee, U.K.; the ⁵Peninsula College of Medicine and Dentistry, University of Exeter, Exeter, U.K.; and the ⁶Trust Centre for Human Genetics, University of Oxford, Oxford, U.K.

Corresponding author: Liling L. Warren, liling.l.warren@gsk.com.

Received 14 July 2011 and accepted 14 January 2012.

DOI: 10.2337/db11-0985

This article contains Supplementary Data online at <http://diabetes.diabetesjournals.org/lookup/suppl/doi:10.2337/db11-0985/-/DC1>.

© 2012 by the American Diabetes Association. Readers may use this article as long as the work is properly cited, the use is educational and not for profit, and the work is not altered. See <http://creativecommons.org/licenses/by-nc-nd/3.0/> for details.

testing corrections were made by adjusting for the total number of tests performed in each study. Meta-analysis was performed using the inverse-variance method (14).

To identify the number of independent SNPs in and around *ADIPOQ*, we conducted variable selection by both frequentist and Bayes methods. In the frequentist approach, both criteria for a SNP to enter or leave the stepwise regression model were set as $P < 0.005$. Bayes variable selection analysis was conducted using the BTAS WinBUGs toolkit (15,16). In both frequentist and Bayes variable selection, missing genotype data were imputed by BEAGLE (17). We further conducted haplotype analysis using Haplo Stats (18) for the set of independent SNPs identified from the aforementioned variable selection analyses to determine whether any of the independent SNPs resided on the same shared haplotype.

In the GO-DARTS study, we tested for association under an additive model using logistic regression for the seven SNPs individually and also in conjunction by analyzing the linear combination of allele dosage data weighted by the effect estimates obtained from linear regression analysis.

Bioinformatics analysis. We used PolyPhen2 (19) and SIFT (20) software to predict impact of nonsynonymous SNPs on the protein function, and PhyloP scores to measure conservation of a base pair across many species (21). Our most strongly associated SNP (rs17366653) is intronic, located 24 nucleotides upstream of the beginning of exon 3. The sequence flanking this position was submitted to a number of splice site or splice branch point bioinformatics prediction tools, including NetGene2 (22), MaxEnt (23), and the Alternative Splicing Desktop (ASD) (<http://www.ebi.ac.uk/asd-srv/wb.cgi?method=6>).

RESULTS

Observed variants. Resequencing of *ADIPOQ* in 14,002 subjects achieved an average depth of 27 reads and an average quality score of 84 (Supplementary Fig. 1). A total of 296 SNPs were observed (Supplementary Table 1). Among them, 52 (18%) were within the coding regions, consisting of 1 nonsense, 30 nonsynonymous, and 21 synonymous variants. Most SNPs were rare, including 169 (57%) singletons and 46 (16%) doubletons, each observed once or twice in 14,002 individuals (Table 1). There were no common nonsynonymous SNPs [minor allele frequency (MAF) $>5\%$] and only two had MAF $> 0.1\%$. The only nonsense variant detected was observed in two European individuals for whom adiponectin measurements were not available.

We compared the frequencies of nonsynonymous SNPs observed in 12,518 European Caucasians, 588 African Americans, and 574 Indian Asians (Supplementary Table 1). Of the 30 nonsynonymous SNPs, 13 were unique to Europeans, 2 to African Americans, and 6 to Indian Asians. Only two, R55C and Y111H, were observed in all three populations. Overall, these variants were extremely rare, and a majority of them were private to each of the three populations included in our study.

Genetic association analyses. We identified three SNPs significantly associated with adiponectin levels. Details of the three SNPs and their effect on adiponectin levels are shown in Table 2. Two of them were previously described,

Y111H (rs17366743) and a 3' untranslated-region SNP (rs6773957). The third (rs17366653), which is the most significantly associated SNP, is previously unreported ($P = 1.02\text{E}-07$, MAF = 0.015 in GEMS; $P = 1.48\text{E}-12$, MAF = 0.020 in CoLaus; meta- $P = 2.20\text{E}-17$). The minor allele was estimated to decrease adiponectin levels by 0.24 $\mu\text{g/mL}$ in the combined samples. Conditional analyses on rs17366743 and rs6773957 yielded highly significant results for rs17366653 ($P < 1.1\text{E}-06$ in GEMS and $P < 2.0\text{E}-09$ in CoLaus), indicating its effect is independent of the other two significant associations in the gene. Four rare nonsynonymous SNPs were found in GEMS and eight in CoLaus. Aggregation analysis gave no interesting results ($P > 0.5$) in GEMS and yielded a modest association in CoLaus ($P = 0.02$) (Supplementary Figs. 2 and 3).

Recently, genome-wide association studies (GWAS) have identified associations between SNPs in and around *ADIPOQ* with adiponectin levels in European populations (7,8,24). For subjects included in our resequencing study with adiponectin measurements, Affymetrix 500 K genotype data were also available. We subsequently analyzed the 353 GWAS SNPs within 2 megabases (Mb) of *ADIPOQ* together with the 15 resequence SNPs with at least 10 copies of the minor allele. Results corresponding to analyses in GEMS, CoLaus, and meta-analysis are shown in Fig. 1. The novel association remained the most statistically significant. The effect of the variant was the largest among all the SNPs being evaluated. Conditional analyses on all other SNPs in this region showed that it was an independent signal from GWAS findings. Furthermore, imputation in a 2-Mb region based on different GWAS panels and data from the 1000 Genomes Project revealed this SNP could not be well imputed ($r^2 < 0.3$). We thus conclude this novel association was previously missed by GWAS and its discovery is due to resequencing a large number of samples.

Seven independent associations (rs17366653, rs17366743, rs1354091, rs3774261, rs3821799, rs16848727, and rs1868146) were identified from the stepwise regression analysis in the GEMS and CoLaus combined dataset. Together, they explained 6% of the adiponectin variation. MAFs of these SNPs ranged from 0.02 to 0.46 (Supplementary Table 2). Bayes variable selection gave very similar results to the frequentist analysis above, with a posterior for the number of independent signals concentrated around 7 and 8 (Supplementary Fig. 4). Haplotype-specific tests corresponded well with individual SNP tests. We thus conclude that these SNPs exert independent effect on adiponectin levels. To examine whether any of the independent SNPs associated with adiponectin levels were contributing to type 2 diabetes risk, we carried out association analyses for the

TABLE 1
Single nucleotide variants observed in *ADIPOQ* in 14,002 individuals by frequency and type

	Nonsense	Nonsynonymous	Synonymous	UTR	Intron	Flanking ⁴	Total
Singleton ¹	0	17	10	131	10	1	169
Doubleton ²	1	5	4	32	2	1	46
(0,0.001] ³	0	6	6	33	1	1	47
(0.001,0.005]	0	1	0	12	0	0	13
(0.005,0.05]	0	1	0	9	1	0	11
(0.05,0.5]	0	0	1	9	0	0	10
Total	1	30	21	226	14	3	296

UTR, untranslated region. ¹Single nucleotide variants observed once in 14,002 individuals. ²Single nucleotide variants observed twice in 14,002 individuals. ³Single nucleotide variants observed more than twice but with minor allele frequencies ≤ 0.001 . ⁴Single nucleotide variants upstream or downstream of gene.

TABLE 2
Characteristics of three significant variants identified in analysis of resequencing data

SNP	Position (NCBI 36.3)	Study	MAF	Minor/major allele	Effect size (SE) (log μ g/mL)	<i>P</i> value
rs17366653	188053510	GEMS	0.015	C/T	−0.460 (0.086)	1.02E−07
rs17366743	188054783	GEMS	0.031	C/T	0.174 (0.064)	6.45E−03
rs6773957	188056399	GEMS	0.391	A/G	0.121 (0.023)	1.70E−07
rs17366653	188053510	CoLaus	0.021	C/T	−0.211 (0.030)	1.48E−12
rs17366743	188054783	CoLaus	0.026	C/T	0.098 (0.027)	2.77E−04
rs6773957	188056399	CoLaus	0.416	A/G	0.035 (0.009)	7.74E−05
rs17366653	188053510	Meta-analysis	0.018	C/T	−0.238 (0.028)	2.20E−17
rs17366743	188054783	Meta-analysis	0.028	C/T	0.109 (0.025)	9.82E−06
rs6773957	188056399	Meta-analysis	0.404	A/G	0.046 (0.008)	2.12E−08

seven SNPs in the GO-DARTS study including 5,145 type 2 diabetic and 6,374 control subjects. None of the tests was statistically significant ($P > 0.1$) (Table 3). However, based on the effect estimate of adiponectin level change on type 2 diabetes risk (1), our statistical power to detect association between these SNPs and type 2 diabetes risk was estimated to range from 6 to 17% for an individual SNP and was at 42% for all seven SNPs combined (Supplementary Table 3).

Testing for association between the seven independent SNPs associated with adiponectin levels and a host of metabolic and cardiovascular-related traits did not yield any significant findings in GEMS and CoLaus (Supplementary Table 4).

Bioinformatics analysis. ASD, MaxEnt, and marginally NetGene2 predicted the minor allele of our most strongly associated SNP (rs17366653) to weaken the scores for splicing at the intron/exon 3 junction. This could lead to aberrant splicing at this site, resulting in an alternative transcript. Further experimental validation is required to validate this hypothesis.

DISCUSSION

By conducting a deep resequencing experiment, we identified a novel association between a low-frequency SNP in

ADIPOQ and adiponectin levels in 3,665 individuals from two independent studies, and confirmed several associations previously identified via GWAS. However, all seven independent SNPs identified only explained about 6% of adiponectin variation in our samples. The unexplained phenotypic variation could be due to structural variations, infrequent noncoding variants, and environmental factors.

Our study provides the most complete assessment of genetic variation in *ADIPOQ* to date. Despite the large number of sequenced subjects, the vast majority of putatively functional variants detected were extremely rare (MAF <0.02%) and likely to be population specific. Recently, a rare (1.1%) nonsynonymous variant G45R of large effect was found in 1,240 Cuban Hispanics (25). This variant is not observed in our samples. The frequency of our most strongly associated SNP was ~1.5% in Europeans, but only 0.3 and 0.6% in African Americans and Indian Asians, respectively. These observations illustrate the limited distribution of such low-frequency variants in different populations and underscore the importance of ethnicity in genetic association studies. An additional challenge of studying very rare variants is the difficulty of replication at the variant level. In CoLaus, we observed a nonsynonymous variant (P91R) of large effect, but only once in 14,002 sequenced subjects.

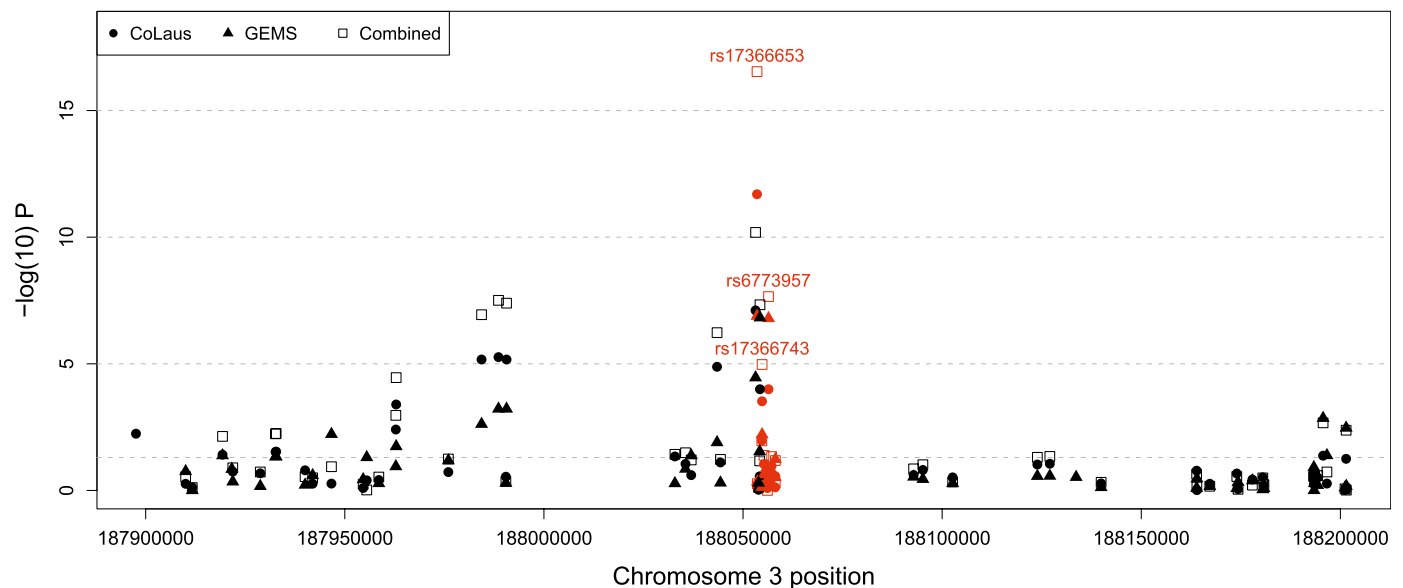


FIG. 1. Association analysis results based on joint analysis of GWAS and resequence SNPs. The red and black symbols represent resequence and GWAS SNPs, respectively.

TABLE 3

Analysis of the seven independent SNPs associated with adiponectin levels in the GO-DARTS study

SNP	<i>P</i> value (adiponectin)	Minor allele	Effect estimate (adiponectin)	<i>P</i> value (type 2 diabetes)	Odds ratio (95% CI) (type 2 diabetes)
rs17366653	2.90E-17	C	-0.24	0.21	1.19 (0.91-1.58)
rs1354091	3.12E-08	G	-0.05	0.21	0.96 (0.90-1.02)
rs3774261	4.66E-08	A	0.04	0.97	1.00 (0.95-1.06)
rs17366743	1.06E-05	C	0.11	0.68	1.03 (0.89-1.21)
rs16848727	0.002	G	0.03	0.11	0.95 (0.89-1.01)
rs1868146	0.05	A	0.03	0.14	1.06 (0.98-1.15)
rs3821799	0.07	T	0.02	0.70	1.01 (0.96-1.07)
Combined				0.43	1.28 (0.69-2.38)

The inverse association between adiponectin levels and type 2 diabetes risk has been well established in epidemiology studies. By focusing on the adiponectin-encoding gene *ADIPOQ*, we were able to rule out concerns over pleiotropic effect of genetic variants associated with other traits. However, all the independent SNPs in *ADIPOQ* only explained 6% of the adiponectin variation in our samples. The estimated statistical power to detect a causal effect of adiponectin on the risk of type 2 diabetes through *ADIPOQ* was only 42% based on our study of 5,145 type 2 diabetic and 6,374 control subjects. Thus our negative finding is inconclusive to tease out the causal relationship between adiponectin and type 2 diabetes. Further investigation by large-scale and well-powered Mendelian randomization studies is warranted.

ACKNOWLEDGMENTS

The CoLaus study was supported by research grants from GlaxoSmithKline and from the Faculty of Biology and Medicine of Lausanne, Switzerland. Funding for the GEMS study was provided by GlaxoSmithKline. Funding for the GO-DARTS study was supported by the Wellcome Trust and genotyping and analysis by Wellcome Trust grant 083270/Z/07/Z. J.R.P. is a Sir Henry Wellcome Postdoctoral Research Fellow (092447/Z/10/Z).

L.L.W., L.L., M.R.N., M.G.E., J.S., D.J.F., J.L.A., K.L.N., A.J.S., P.M.W., M.D.H., S.D.T., X.Y., L.R.C., S.L.C., V.M., J.C.W., and D.M.W. are employees of GlaxoSmithKline. No other potential conflicts of interest relevant to this article were reported.

L.L.W. researched data and wrote the manuscript. L.L., J.S., D.J.F., J.L.A., K.L.N., A.J.S., and X.Y. researched data. M.R.N., M.G.E., S.D.T., S.L.C., T.M.F., J.C.W., and D.M.W. contributed to discussion and reviewed and edited the manuscript. P.M.W., M.D.H., L.R.C., V.M., A.D.M., C.N.A.P., and J.R.P. reviewed and edited the manuscript. L.L.W. is the guarantor of this work and, as such, had full access to all the data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis.

The authors thank the co-primary investigators of the CoLaus study, Gerard Waeber (Centre Hospitalier Universitaire Vaudois, Lausanne, Switzerland) and Peter Vollenweider (Centre Hospitalier Universitaire Vaudois), and the primary investigator of the PsyColaus Study, Martin Preisig (Centre Hospitalier Universitaire Vaudois). The authors thank GEMS investigators: Scott Grundy (UT Southwestern Medical Center, Dallas, Texas), Jonathan Cohen (UT Southwestern Medical Center), Ruth McPherson (Ottawa Heart Institute, Ottawa, Canada), Antero Kesaniemi (University of Oulu, Oulu, Finland), Robert Mahley (University

of California, San Francisco, CA), Tom Bersot (University of California, San Francisco, CA), Philip Barter (The University of Adelaide, Adelaide, Australia), and Gerard Waeber (Centre Hospitalier Universitaire Vaudois). The authors are grateful to John Novembre and Daniel Wegmann (University of California, Los Angeles, CA) for helpful discussions. The authors thank study participants who provided written informed consent for the use of their DNA in genetic studies.

REFERENCES

- Li S, Shin HJ, Ding EL, van Dam RM. Adiponectin levels and risk of type 2 diabetes: a systematic review and meta-analysis. *JAMA* 2009;302:179-188
- Ebrahim S, Davey Smith G. Mendelian randomization: can genetic epidemiology help redress the failures of observational epidemiology? *Hum Genet* 2008;123:15-33
- Smith GD, Ebrahim S. Mendelian randomization: prospects, potentials, and limitations. *Int J Epidemiol* 2004;33:30-42
- Hivert MF, Manning AK, McAteer JB, et al. Common variants in the adiponectin gene (*ADIPOQ*) associated with plasma adiponectin levels, type 2 diabetes, and diabetes-related quantitative traits: the Framingham Offspring Study. *Diabetes* 2008;57:3353-3359
- Jee SH, Sull JW, Lee JE, et al. Adiponectin concentrations: a genome-wide association study. *Am J Hum Genet* 2010;87:545-552
- Lindsay RS, Funahashi T, Krakoff J, et al. Genome-wide linkage analysis of serum adiponectin in the Pima Indian population. *Diabetes* 2003;52:2419-2425
- Ling H, Waterworth DM, Stirradd HA, et al. Genome-wide linkage and association analyses to identify genes influencing adiponectin levels: the GEMS study. *Obesity (Silver Spring)* 2009;17:737-744
- Richards JB, Waterworth D, O'Rahilly S, et al.; GIANT Consortium. A genome-wide association study reveals variants in *ARL15* that influence adiponectin levels. *PLoS Genet* 2009;5:e1000768
- Stirradd H, Lin X, Ling H, et al. Genetic and phenotypic architecture of metabolic syndrome-associated components in dyslipidemic and normolipidemic subjects: the GEMS study. *Atherosclerosis* 2008;197:868-876
- Firmann M, Mayor V, Vidal PM, et al. The CoLaus study: a population-based study to investigate the epidemiology and genetic determinants of cardiovascular risk factors and metabolic syndrome. *BMC Cardiovasc Disord* 2008;8:6
- Zeggini E, Scott LJ, Saxena R, et al.; Wellcome Trust Case Control Consortium. Meta-analysis of genome-wide association data and large-scale replication identifies additional susceptibility loci for type 2 diabetes. *Nat Genet* 2008;40:638-645
- Li R, Li Y, Fang X, et al. SNP detection for massively parallel whole-genome resequencing. *Genome Res* 2009;19:1124-1132
- Li B, Leal SM. Methods for detecting associations with rare variants for common diseases: application to analysis of sequence data. *Am J Hum Genet* 2008;83:311-321
- DerSimonian R, Laird N. Meta-analysis in clinical trials. *Control Clin Trials* 1986;7:177-188
- Lunn DJ, Whittaker JC, Best N. A Bayesian toolkit for genetic association studies. *Genet Epidemiol* 2006;30:231-247
- Lunn D, Thomas A, Best N, Spiegelhalter D. WinBUGS-a Bayesian modelling framework: concepts, structure, and extensibility. *Stat Comput* 2000;10:325-337
- Browning SR, Browning BL. Rapid and accurate haplotype phasing and missing-data inference for whole-genome association studies by use of localized haplotype clustering. *Am J Hum Genet* 2007;81:1084-1097

18. Schaid DJ, Rowland CM, Tines DE, Jacobson RM, Poland GA. Score tests for association between traits and haplotypes when linkage phase is ambiguous. *Am J Hum Genet* 2002;70:425–434
19. Ramensky V, Bork P, Sunyaev S. Human non-synonymous SNPs: server and survey. *Nucleic Acids Res* 2002;30:3894–3900
20. Ng PC, Henikoff S. Predicting deleterious amino acid substitutions. *Genome Res* 2001;11:863–874
21. Siepel A, Bejerano G, Pedersen JS, et al. Evolutionarily conserved elements in vertebrate, insect, worm, and yeast genomes. *Genome Res* 2005;15:1034–1050
22. Brunak S, Engelbrecht J, Knudsen S. Prediction of human mRNA donor and acceptor sites from the DNA sequence. *J Mol Biol* 1991;220:49–65
23. Desmet FO, Hamroun D, Lalande M, Collod-Bérout G, Claustres M, Bérout C. Human Splicing Finder: an online bioinformatics tool to predict splicing signals. *Nucleic Acids Res* 2009;37:e67
24. Heid IM, Wagner SA, Gohlke H, et al. Genetic architecture of the APM1 gene and its influence on adiponectin plasma levels and parameters of the metabolic syndrome in 1,727 healthy Caucasians. *Diabetes* 2006;55:375–384
25. Bowden DW, An SS, Palmer ND, et al. Molecular basis of a linkage peak: exome sequencing and family-based analysis identify a rare genetic variant in the ADIPOQ gene in the IRAS family study. *Hum Mol Genet* 2010;19:4112–4120